

Announced Care Inspection Report 18 October 2018



Ardent Dental Care

Type of Service: Independent Hospital (IH) – Dental Treatment

Address: 14 Portaferry Road, Newtownards BT23 8NN

Tel No: 028 9182 1348

Inspector: Norma Munn

www.rqia.org.uk

Assurance, Challenge and Improvement in Health and Social Care

It should be noted that this inspection report should not be regarded as a comprehensive review of all strengths and areas for improvement that exist in the service. The findings reported on are those which came to the attention of RQIA during the course of this inspection. The findings contained within this report do not exempt the service from their responsibility for maintaining compliance with legislation, standards and best practice.

1.0 What we look for



In respect of dental practices for the 2018/19 inspection year we are moving to a more focused, shorter inspection which will concentrate on the following key patient safety areas:

- management of medical emergencies
- infection prevention and control
- decontamination of reusable dental instruments
- radiology and radiation safety
- review of areas for improvement from the last inspection

2.0 Profile of service

This is a registered dental practice with three registered places.

3.0 Service details

Organisation/Registered Provider: Mr Simon Gay	Registered Manager: Mr Simon Gay
Person in charge at the time of inspection: Mr Simon Gay	Date manager registered: 6 March 2012
Categories of care: Independent Hospital (IH) – Dental Treatment	Number of registered places: 3

4.0 Action/enforcement taken following the most recent inspection dated 29 March 2018

The most recent inspection of the establishment was an announced care inspection. The completed quality improvement plan (QIP) was returned and approved by the care inspector.

4.1 Review of areas for improvement from the last care inspection dated 29 March 2018

Areas for improvement from the last care inspection		
Action required to ensure compliance with The Minimum Standards for Dental Care and Treatment (2011)		Validation of compliance
Area for improvement 1 Ref: Standard 11 Stated: Second time	A system should be implemented for appraising staff performance at least on an annual basis.	Met
	Action taken as confirmed during the inspection: Discussion with Mr Gay and a review of documentation confirmed that a system has been implemented for appraising staff performance and Mr Gay confirmed that appraisals would take place on an annual basis.	

5.0 Inspection findings

An announced inspection took place on 18 October 2018 from 09.50 to 11.20.

This inspection was underpinned by The Health and Personal Social Services (Quality, Improvement and Regulation) (Northern Ireland) Order 2003, The Independent Health Care Regulations (Northern Ireland) 2005, The Regulation and Improvement Authority (Independent Health Care) (Fees and Frequency of Inspections) (Amendment) Regulations (Northern Ireland) 2011 and the Department of Health (DOH) Minimum Standards for Dental Care and Treatment (2011).

A poster informing patients that an inspection was being conducted was displayed.

During the inspection the inspector met with Mr Gay, registered person, one associate dentist, two dental nurses and two receptionists. A tour of some areas of the premises was also undertaken.

The findings of the inspection were provided to Mr Gay at the conclusion of the inspection.

5.1 Management of medical emergencies

Management of medical emergencies

A review of arrangements in respect of the management of a medical emergency evidenced that, in the main, emergency medicines were provided in keeping with the British National Formulary (BNF). It was observed that Adrenaline was retained in an autoinjector format in two doses. Adrenaline should be available in three doses, which are, 150 micrograms, 300 micrograms and 500 micrograms with sufficient stock to be able to administer a second dose to the same patient if necessary in keeping with the BNF and the Health and Social Care Board (HSCB) guidance. A discussion took place in regards to the procedure for the safe administration of Adrenaline and the various doses and quantities as recommended by the HSCB and BNF and Mr Gay confirmed that he had ordered a supply of Adrenaline ampoules prior to the inspection and was awaiting delivery of these. Following the inspection RQIA received confirmation that Adrenaline ampoules had been provided. The Glucagon medication had been stored in the fridge and fridge temperatures had not been recorded. Mr Gay was advised that if Glucagon is stored in the fridge, daily fridge temperatures should be taken and recorded to evidence that the cold chain has been maintained. Mr Gay was advised that as there would be no evidence that the cold chain has been maintained a revised expiry date of 18 months from the date of receipt should be marked on the medication packaging and expiry date checklist to reflect that the cold chain has been broken as per manufacturer's instructions. This was actioned on the day of the inspection.

Emergency equipment as recommended by the Resuscitation Council (UK) guidelines was retained. A system was in place to ensure that emergency medicines and equipment do not exceed their expiry date.

Review of training records and discussion with staff confirmed that the management of medical emergencies is included in the induction programme and training is updated on an annual basis in keeping with best practice guidance. The most recent occasion staff completed medical emergency refresher training was during November 2017.

Discussion with staff demonstrated that they have a good understanding of the actions to be taken in the event of a medical emergency and the location of medical emergency medicines and equipment.

Areas of good practice

The review of the arrangements in respect of the management of a medical emergency confirmed that this dental practice takes a proactive approach to this key patient safety area. This includes ensuring that staff have the knowledge and skills to react to a medical emergency, should it arise.

Areas for improvement

No further areas for improvement were identified during the inspection.

	Regulations	Standards
Areas for improvement	0	0

5.2 Infection prevention and control

Infection prevention and control (IPC)

During a tour of the premises, it was evident that the practice, including the clinical and decontamination areas, was clean, tidy and uncluttered.

The practice continues to audit compliance with Health Technical Memorandum (HTM) 01-05: Decontamination in primary care dental practices using the Infection Prevention Society (IPS) audit tool. This audit includes key elements of IPC, relevant to dentistry, including the arrangements for environmental cleaning, the use of personal protective equipment, hand hygiene practice, and waste and sharps management.

A review of the most recent IPS audit, completed during March/April 2018, evidenced that the audit had been completed in a meaningful manner and had identified areas of good practice. Mr Gay confirmed that any learning identified as a result of these audits is shared with staff when identified. Following the inspection RQIA received confirmation that the IPS audit for October 2018 had been undertaken.

Arrangements were in place to ensure that staff received IPC training commensurate with their roles and responsibilities and during discussion with staff it was confirmed that they had a good level of knowledge and understanding of IPC procedures.

During discussion it was identified that conventional needles and syringes are used by the dentists when administering local anaesthetic as opposed to using safer sharps. Regulation 5 (1) (b) of The Health and Safety (Sharp Instruments in Healthcare) Regulations (Northern Ireland) 2013 states that ‘safer sharps are used so far as is reasonably practicable’. Mr Gay confirmed that it is the responsibility of the user of sharps to safely dispose of them. Sharps risk assessments were not in place for the dentists who do not use safer sharps. Mr Gay was advised that safer sharps should be used so far as is reasonably practicable, in keeping with legislation, and where this is not practicable a risk assessment should be completed in respect of each dentist. Following the inspection RQIA received confirmation that a risk assessment had been completed on the management of sharps and a copy of the risk assessment had been shared with the other dentists in the practice.

Areas of good practice

A review of the current arrangements evidenced that standards in respect of infection prevention and control practice are being given high priority. This includes proactively auditing practice, taking action when issues are identified and ensuring staff have the knowledge and skills to ensure standards are maintained.

Areas for improvement

No further areas for improvement were identified during the inspection.

	Regulations	Standards
Areas for improvement	0	0

5.3 Decontamination of reusable dental instruments

Decontamination of reusable dental instruments

A decontamination room separate from patient treatment areas and dedicated to the decontamination process was available. The decontamination room facilitates the flow from dirty through to clean areas for the cleaning and sterilising of reusable instruments.

The processes in respect of the decontamination of reusable dental instruments are being audited in line with best practice outlined in HTM 01-05 using the IPS audit tool.

As discussed a review of the most recent IPS audit, completed during March/April 2018, evidenced that the audit had been completed in a meaningful manner and had identified areas of good practice.

Arrangements were in place to ensure that staff receive training in respect of the decontamination of reusable dental instruments commensurate with their roles and responsibilities.

A review of current practice evidenced that arrangements are in place to ensure that reusable dental instruments are appropriately cleaned, sterilised and stored following use in keeping with best practice guidance as outlined in HTM 01-05.

Appropriate equipment, including a washer disinfector, two DAC Universals and one steam steriliser, has been provided to meet the practice requirements. Mr Gay confirmed that the steriliser has been decommissioned and his intention is to purchase a new steriliser in the future. On the day of the inspection one of the DAC Universals and the washer disinfector were out of action. However, Mr Gay confirmed that this had only been for a short period of time and immediately following the inspection RQIA received confirmation that washer disinfector had been fixed and the practice are waiting for a part to be delivered in respect of the DAC Universal.

The new DAC Universal had been validated during March 2018. Mr Gay confirmed that a date had been arranged for the validation of the washer disinfector and the second DAC Universal. Following the inspection RQIA received confirmation that this equipment had been validated on 1 November 2018.

The decontamination equipment had been inspected in keeping with the written scheme of examination and equipment logbooks evidenced that periodic tests are undertaken and recorded in keeping with HTM 01-05.

Staff are aware of what equipment in the practice should be treated as single use and what equipment is suitable for decontamination. It was confirmed that single use devices are only used for single-treatment episodes and disposed of following use.

Areas of good practice

A review of the current arrangements evidenced that best practice as outlined in HTM 01-05 is being achieved in respect of the decontamination of reusable dental instruments. This includes proactively auditing practice, taking action when issues are identified and ensuring staff have the knowledge and skills to ensure standards are maintained.

Areas for improvement

No areas for improvement were identified during the inspection.

	Regulations	Standards
Areas for improvement	0	0

5.4 Radiology and radiation safety

Radiology and radiation safety

The practice has three surgeries, each of which has an intra-oral x-ray machine.

Mr Gay is the radiation protection supervisor (RPS) for the practice. Mr Gay was aware of the most recent changes to the legislation surrounding radiology and radiation safety and a radiation protection advisor (RPA) and medical physics expert (MPE) have been appointed.

A dedicated radiation protection file containing all relevant information was in place. Mr Gay confirmed that he regularly reviews the information contained within the file to ensure that it is current.

The appointed RPA completes a quality assurance check every three years. A review of the report of the most recent visit by the RPA demonstrated that any recommendations made have been addressed.

Staff spoken with demonstrated sound knowledge of radiology and radiation safety in keeping with their roles and responsibilities.

Mr Gay takes a proactive approach to radiation safety and protection by conducting a range of audits, including x-ray quality grading and justification and clinical evaluation recording.

Areas of good practice

A review of radiology and radiation safety arrangements evidenced that the radiation protection supervisor for this practice takes a proactive approach to the management of radiology and radiation safety.

Areas for improvement

No areas for improvement were identified during the inspection.

	Regulations	Standards
Areas for improvement	0	0

5.5 Equality data

Equality data

The arrangements in place in relation to the equality of opportunity for patients and the importance of staff being aware of equality legislation and recognising and responding to the diverse needs of patients was discussed with Mr Gay and staff.

5.6 Patient and staff views

Eight patients submitted questionnaire responses to RQIA. All of the patients indicated that they felt their care was safe and effective, that they were treated with compassion and that the service was well led and were very satisfied with each of these areas of their care. One comment was included in a submitted questionnaire response as follows:

- “I am always very pleased with the standard of care and treatment I receive.”

RQIA invited staff to complete an electronic questionnaire prior to the inspection. No staff submitted questionnaire responses to RQIA.

5.7 Total number of areas for improvement

	Regulations	Standards
Total number of areas for improvement	0	0

6.0 Quality improvement plan

There were no areas for improvement identified during this inspection, and a QIP is not required or included, as part of this inspection report.



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